Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	2	("7153984").PN.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2007/06/12 06:00
L2	3	("3920582").PN.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2007/06/12 06:01
L3		("3053884").PN.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2007/06/12 06:02
L4	0	("methanetrisulfonic").PN.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2007/06/12 07:40
`L5	. 18	methanetrisulfonic	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 08:32
L6	374665	phenol	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 06:03
L7	5	IS and I6	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 06:03
L8	5	IS same I6	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 06:03
L9	0	("3053884").URPN.	USPAT	OR	ON	2007/06/12 06:09
L10	2	("6103924").PN.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2007/06/12 06:10
L11	2	("20020004619").PN.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2007/06/12 06:38
L12	1	("4247720").PN.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2007/06/12 06:43

L13	126	ketoisophorone	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 06:43
L14	1	I5 and I13	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 06:43
L15	213883	sulfonic	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 06:44
L16	22	I13 and I15	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 06:44
L18	1	I13 same I15	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 06:48
L19	. 17	"2149159"	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 07:03
L20	2	("3082258").PN.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2007/06/12 07:09
L21	1	"19805690"	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 07:26
L22	580	560/254.ccls.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 07:39
L23	0	l13 and l22	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 07:27
L24	0	I5 and I22	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 07:27
L25	64920	hydroquinone	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 07:29

		1	-	·	,	·
L26	35	I22 and I25	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 07:29
L27	67055	\$hydroquinone	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 07:39
L28	0	("methanesulfonic").PN.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2007/06/12 07:40
L29	52312	methanesulfonic	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 07:40
L30	174	methanedisulfonic	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 07:55
L31	0	l13 and l30	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 07:41
L32	9	l13 and l29	US-PGPUB; USPAT; EPO; JPO; DERWENT	ÖR	ON	2007/06/12 07:41
L33	0	(methanetrisulfonic and (ketoisophorone or "3,5,5-trimethyl-l, 4-benzoquinone") and hydroquinone). clm.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON .	2007/06/12 08:32
L34	0	(methanetrisulfonic and (ketoisophorone or "3,5,5-trimethyl-I, 4-benzoquinone")).clm.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 08:33
L36	2	("20070123720").PN.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2007/06/12 08:31
L37	1	(methane adj trisulfonic and (ketoisophorone or "3,5,5-trimethyl-l, 4-benzoquinone") and hydroquinone). clm.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 08:32
L38	19	methane adj trisulfonic	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 08:32

L39	1	(methane adj trisulfonic and (ketoisophorone or "3,5,5-trimethyl-l, 4-benzoquinone")).clm.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 08:33
L40	17	138 not 15	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 08:34
L41	16	I5 not I38	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/06/12 08:42

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     1
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NEWS
      2
         JAN 08
NEWS
      3
         JAN 16
                 CA/CAplus Company Name Thesaurus enhanced and reloaded
NEWS
     4
         JAN 16
                 IPC version 2007.01 thesaurus available on STN
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         JAN 16
                 WPIDS/WPINDEX/WPIX enhanced with IPC 8 reclassification data
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     6 JAN 22
                 CA/CAplus updated with revised CAS roles
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                 CA/CAplus enhanced with patent applications from India
NEWS 8 JAN 29
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NEWS 9 JAN 29
                 CAS Registry Number crossover limit increased to 300,000 in
                 multiple databases
NEWS 10 FEB 15
                 PATDPASPC enhanced with Drug Approval numbers
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        FEB 15
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                KOREAPAT enhanced with IPC 8 features and functionality
NEWS 13 FEB 26
                MEDLINE reloaded with enhancements
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                EMBASE enhanced with Clinical Trial Number field
NEWS 15 FEB 26
                TOXCENTER enhanced with reloaded MEDLINE
NEWS 16 FEB 26
                IFICDB/IFIPAT/IFIUDB reloaded with enhancements
NEWS 17 FEB 26
                CAS Registry Number crossover limit increased from 10,000
                 to 300,000 in multiple databases
NEWS 18
        MAR 15
                WPIDS/WPIX enhanced with new FRAGHITSTR display format
NEWS 19 MAR 16
                CASREACT coverage extended
NEWS 20 MAR 20
                MARPAT now updated daily
NEWS 21 MAR 22 LWPI reloaded
NEWS 22 MAR 30
                RDISCLOSURE reloaded with enhancements
NEWS 23 APR 02
                JICST-EPLUS removed from database clusters and STN
NEWS 24 APR 30
                GENBANK reloaded and enhanced with Genome Project ID field
NEWS 25 APR 30
                CHEMCATS enhanced with 1.2 million new records
NEWS 26 APR 30
                CA/CAplus enhanced with 1870-1889 U.S. patent records
NEWS 27 APR 30
                 INPADOC replaced by INPADOCDB on STN
NEWS 28 MAY 01
                New CAS web site launched
NEWS 29
        MAY 08
                 CA/CAplus Indian patent publication number format defined
NEWS 30
        MAY 14
                RDISCLOSURE on STN Easy enhanced with new search and display
                 fields
NEWS 31
        MAY 21
                BIOSIS reloaded and enhanced with archival data
NEWS 32
        MAY 21
                TOXCENTER enhanced with BIOSIS reload
NEWS 33
        MAY 21
                CA/CAplus enhanced with additional kind codes for German
                patents
NEWS 34
        MAY 22
                CA/CAplus enhanced with IPC reclassification in Japanese
                patents
NEWS EXPRESS NOVEMBER 10 CURRENT WINDOWS VERSION IS V8.01c, CURRENT
             MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
             AND CURRENT DISCOVER FILE IS DATED 25 SEPTEMBER 2006.
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=> e	ketoisophoron	e/cn
E1	1	KETOISDIN/CN
E2	1	KETOISOLACTARORUFIN/CN
E3	1>	KETOISOPHORONE/CN
E4	1	KETOISOSTEVIC ACID/CN
E5	1	KETOISOVALERATE OXIDOREDUCTASE SUBUNIT (METHANOSARCINA BARKE
77.6	•	RI STRAIN FUSARO)/CN
E6	1	KETOISOVALERATE OXIDOREDUCTASE SUBUNIT (METHANOSARCINA MAZEI STRAIN GOEl GENE VORA)/CN
E7	1	KETOISOVALERATE OXIDOREDUCTASE SUBUNIT (METHANOSARCINA MAZEI STRAIN GOE1 GENE VORB)/CN
E8	1	KETOISOVALERATE OXIDOREDUCTASE SUBUNIT (METHANOSARCINA MAZEI
50	-	STRAIN GOE1 GENE VORC)/CN
E9	2	KETOISOVALERATE OXIDOREDUCTASE SUBUNIT VORA (BACTEROIDES FRA
		GILIS STRAIN YCH46)/CN
E10	1	KETOISOVALERATE OXIDOREDUCTASE SUBUNIT VORA (BACTEROIDES THE
		TAIOTAOMICRON STRAIN VPI-5482 GENE BT0329)/CN
E11	1	KETOISOVALERATE OXIDOREDUCTASE SUBUNIT VORA (BACTEROIDES THE
		TAIOTAOMICRON STRAIN VPI-5482 GENE BT0330)/CN
E12	1	KETOISOVALERATE OXIDOREDUCTASE SUBUNIT VORB (BACTEROIDES FRA
		GILIS STRAIN YCH46)/CN

=> d l1

L1 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2007 ACS on STN RN 1125-21-9 REGISTRY ED Entered STN: 16 Nov 1984 2-Cyclohexene-1,4-dione, 2,6,6-trimethyl- (CA INDEX NAME) CN OTHER NAMES: CN 2,6,6-Trimethyl-2-cyclohexen-1,4-dione CN 2,6,6-Trimethylcyclohex-2-ene-1,4-dione CN 3,5,5-Trimethyl-2-cyclohexene-1,4-dione CN 4-Ketoisophorone CN $4-0xo-\alpha$ -isophorone 4-Oxoisophorone CN 6-Oxoisophorone CN keto-Isophorone CN Ketoisophorone CN CN Oxoisophorone CN Oxopholone CN Oxophorone MF C9 H12 O2 AGRICOLA, ANABSTR, BEILSTEIN*, BIOSIS, CA, CAOLD, CAPLUS, LC STN Files: CASREACT, CHEMCATS, CHEMINFORMRX, CHEMLIST, CSCHEM, IFICDB, IFIPAT, IFIUDB, MEDLINE, SPECINFO, TOXCENTER, USPAT2, USPATFULL, VTB (*File contains numerically searchable property data) DSL**, EINECS**, TSCA** Other Sources: (**Enter CHEMLIST File for up-to-date regulatory information)

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390 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA 396 REFERENCES IN FILE CAPLUS (1907 TO DATE) 7 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

TOTAL

7.56

=> file caplus COST IN U.S. DOLLARS SINCE FILE ENTRY SESSION FULL ESTIMATED COST 7.35

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FULL ESTIMATED COST
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8.03

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http://www.cas.org/support/stngen/stndoc/properties.html

```
=> e metrhanetrisulfonic acid/cn
E1
            2
                 METRETON/CN ·
E2
            1
                  METREX/CN
            0 --> METRHANETRISULFONIC ACID/CN
E3
E4
            1
                 METRIAN/CN
E5
            1
                 METRIAR/CN
E6
            1
                 METRIAREZ-B/CN
E7
            1
                 METRIAREZ-Γ/CN
E8
           1
                 METRIBEN/CN
E9
           1
                 METRIBOLONE/CN
E10
            1
                 METRIBUSIN-BENEFIN MIXTURE/CN
           . 1
E11
                 METRIBUZIN/CN
                METRIBUZIN DA/CN
E12
=> e methanetrisulfonic acid/cn
                METHANETRISULFONAMIDE, N, N', N''-TRIS(1-PHENYLETHYL)-/CN
El
            1
                  METHANETRISULFONAMIDE, N,N',N''-TRIS(TRIMETHYLSILYL)-/CN
E2
E3
            1 --> METHANETRISULFONIC ACID/CN
E4
            1 METHANETRISULFONIC ACID, ALUMINUM SALT/CN
E5
                  METHANETRISULFONIC ACID, BARIUM SALT (2:3)/CN
            1
E6
                  METHANETRISULFONIC ACID, BROMO-/CN
```

```
METHANETRISULFONIC ACID, BROMO-, TRIETHYL ESTER/CN
E7
             1
                   METHANETRISULFONIC ACID, BROMO-, TRIMETHYL ESTER/CN
E8
             1
                   METHANETRISULFONIC ACID, BROMO-, TRIS(TRIMETHYLSILYL) ESTER/
E9
             1
                   CN
E10
                   METHANETRISULFONIC ACID, CHLORO-/CN
             1
                   METHANETRISULFONIC ACID, CHLORO-, TRIS(TRIMETHYLSILYL) ESTER
E11
             1
                   METHANETRISULFONIC ACID, FLUORO-/CN
E12
             1
=> e3
             1 "METHANETRISULFONIC ACID"/CN
1.2
=> d 12
     ANSWER 1 OF 1 REGISTRY COPYRIGHT 2007 ACS on STN
L2
RN
     54322-33-7 REGISTRY
ED
    Entered STN: 16 Nov 1984
    Methanetrisulfonic acid (7CI, '9CI) (CA INDEX NAME)
CN
DR
     856207-29-9
    C H4 O9 S3
MF
CI
    COM
LC
     STN Files: BEILSTEIN*, CA, CAOLD, CAPLUS, CASREACT, CHEMLIST, CSCHEM,
       GMELIN*, IFICDB, IFIPAT, IFIUDB, USPAT2, USPATFULL
         (*File contains numerically searchable property data)
     Other Sources:
                      EINECS**
         (**Enter CHEMLIST File for up-to-date regulatory information)
     SO<sub>3</sub>H
HO3S-CH-SO3H
**PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**
              25 REFERENCES IN FILE CA (1907 TO DATE)
               2 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
              25 REFERENCES IN FILE CAPLUS (1907 TO DATE)
               2 REFERENCES IN FILE CAOLD (PRIOR TO 1967)
=> e sulfonic acid/cn
E1
             1
                   SULFONIAZIDE NITRATE/CN
E2
                   SULFONIAZIDE SODIUM/CN
E3
             0 --> SULFONIC ACID/CN
E4
             1
                  SULFONIC ACID LS/CN
E5
             1
                   SULFONIC ACID, ((3,5-BIS(1,1-DIMETHYLETHYL)-4-HYDROXYPHENYL)
                   METHYL) - , MONOBUTYL ESTER, NICKEL COMPLEX/CN
E6
             1
                   SULFONIC ACID, PHOSPHINO-/CN
E7
                   SULFONIC ACIDS/CN
E8
                  SULFONIC ACIDS, ALKANE, CHLORO/CN
E9
                  SULFONIC ACIDS, ALKANE, CHLORO, SODIUM SALTS/CN
E10
                 SULFONIC ACIDS, ALKANE, SODIUM SALTS/CN
E11
                 SULFONIC ACIDS, ALKANEDI-, DISODIUM SALTS/CN
E12
                  SULFONIC ACIDS, ALKANESULFONIC, CHLORO/CN
=> e methanesulfonic acid/cn
El
                   METHANESULFONATE SULFONATASE MSUD (PSEUDOMONAS FLUORESCENS S
                   TRAIN PF-5 GENE MSUD)/CN
                  METHANESULFONATE SULFONATASE; MSUD (MESORHIZOBIUM LOTI STRAI
E2
                   N MAFF303099 GENE MLR5216)/CN
E3
             1 --> METHANESULFONIC ACID/CN
E4
             1
                   METHANESULFONIC ACID ((3'-((5-((TERT-BUTOXYCARBONYLAMINO)(IM
                   INO) METHYL) -2- (METHYLSULFANYL) THIEN-3-YL) SULFONYL) -4- (N, N'-B
```

```
IS (TERT-BUTOXYCARBONYL) GUANIDINO) - 6 - METHYLBIPHENYL - 2 - YL) CARB
                    AMOYL) METHYL ESTER/CN
                   METHANESULFONIC ACID ((3S,4R)-4-(((5-CHLOROTHIEN-2-YL)CARBON
E5
                    YL) AMINO) -1-(((2-FLUORO-4-(2-OXO-2H-PYRIDIN-1-YL) PHENYL) CARB
                    AMOYL) METHYL) PYRROLIDIN-3-YL) METHYL ESTER/CN ·
                    METHANESULFONIC ACID ((5R)-3-(3-FLUORO-4-(TETRAHYDROTHIOPYRA
E6
                    N-4-YL) PHENYL) -2-OXOOXAZOLIDIN-5-YL) METHYL ESTER/CN
                   METHANESULFONIC ACID ((5R)-3-(4-((1,4-DIBENZYLPIPERAZIN-2-YL
E7
                   METHYL) ETHYLAMINO) - 3 - FLUOROPHENYL) - 2 - OXOOXAZOLIDIN - 5 - YL) METH
                    YL ESTER/CN
                   METHANESULFONIC ACID ((5R)-3-(4-(3,6-DIHYDRO-2H-THIOPYRAN-4-
E8
                    YL) -3,5-DIFLUOROPHENYL) -2-OXOOXAZOLIDIN-5-YL) METHYL ESTER/CN
                   METHANESULFONIC ACID ((5R)-3-(4-(3,6-DIHYDRO-2H-THIOPYRAN-4-
E9
             1
                    YL) - 3 - FLUOROPHENYL) - 2 - OXOOXAZOLIDIN - 5 - YL) METHYL ESTER/CN
                    METHANESULFONIC ACID ((R)-2,2-DIMETHYL-(1,3)DIOXOLAN-4-YL)ME
E10
             1
                    THYL ESTER/CN
                   METHANESULFONIC ACID ((R)-2-OXO-3-(1-OXO-3-(2-TRIFLUOROMETHY
E11
             1
                    LPHENYL) -1.2-DIHYDROISOOUINOLIN-7-YL) OXAZOLIDIN-5-YL) METHYL
                    ESTER/CN
                   METHANESULFONIC ACID ((R)-2-OXO-3-(8-OXO-6,7,8,9-TETRAHYDRO-
E12
             1
                    5H-BENZOCYCLOHEPTEN-2-YL)OXAZOLIDIN-5-YL)METHYL ESTER/CN
=> e3
             1 "METHANESULFONIC ACID"/CN
L3
=> d 13
     ANSWER 1 OF 1 REGISTRY COPYRIGHT 2007 ACS on STN
L3
RN
     75-75-2 REGISTRY
ED
     Entered STN: 16 Nov 1984
     Methanesulfonic acid (CA INDEX NAME)
CN
OTHER NAMES:
     MCAT 1201
CN
CN
     Methylsulfonic acid
CN
     NSC 3718
     125756-91-4, 98527-29-8, 115449-98-4, 62203-24-1, 87128-90-3, 44209-64-5,
DR
     44209-72-5
MF
     C H4 O3 S
CI
     COM
LC
     STN Files:
                   AGRICOLA, ANABSTR, BEILSTEIN*, BIOSIS, BIOTECHNO, CA, CAOLD,
       CAPLUS, CASREACT, CBNB, CHEMCATS, CHEMINFORMRX, CHEMLIST, CIN, CSCHEM,
       DETHERM*, EMBASE, ENCOMPLIT, ENCOMPLIT2, ENCOMPPAT, ENCOMPPAT2, GMELIN*,
       HSDB*, IFICDB, IFIPAT, IFIUDB, IPA, MEDLINE, MRCK*, MSDS-OHS, PIRA,
       PROMT, PS, RTECS*, SPECINFO, SYNTHLINE, TOXCENTER, ULIDAT, USPAT2,
       USPATFULL, VTB
         (*File contains numerically searchable property data)
                      DSL**, EINECS**, TSCA**
         (**Enter CHEMLIST File for up-to-date regulatory information)
```

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5744 REFERENCES IN FILE CA (1907 TO DATE)
205 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
5765 REFERENCES IN FILE CAPLUS (1907 TO DATE)
21 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

=> file caplus
COST IN U.S. DOLLARS

FULL ESTIMATED COST

SINCE FILE TOTAL ENTRY SESSION 15.15 23.18

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=> 11 and 12

396 L1

25 L2

L4 0 L1 AND L2

=> l1

L5 396 L1

=> 12

L6 25 L2

=> 13

L7 5760 L3

=> d 16 1-25 ti

- L6 ANSWER 1 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Acylation process for the preparation of 2,3,5-trimethylhydroquinone diacylates in the presence of methanetrisulfonic acid catalyst
- L6 ANSWER 2 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Process for the manufacture of tocyl and tocopheryl acylates
- L6 ANSWER 3 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Crosslinked polyoxyalkylene-polysiloxanes for use as nonaqueous salt-type electrolytes for lithium secondary batteries
- L6 ANSWER 4 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Photochemical crosslinked epoxy-polyoxyalkylene-polysiloxanes for use as nonaqueous salt-type electrolytes for lithium secondary batteries
- L6 ANSWER 5 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Manufacture of α -tocopherol from the reaction of trimethylhydroquinone with isophytol or phytol in the presence of methane trisulfonate

- L6 ANSWER 6 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Investigations on the acid-strength of alkanepolysulfonic acids
- L6 ANSWER 7 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Improvement of the synthesis of the tripotassium methanetrisulfonate monohydrate
- L6 ANSWER 8 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Process for the preparation of alkyl 3-alkoxypropionates
- L6 ANSWER 9 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI The electrochemistry of a dimeric and two monomeric cistrioxomolybdenum(VI) complexes containing cyclic triamine ligands in protic and aprotic media: model compounds for the active site in formate dehydrogenase
- L6 ANSWER 10 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Alkoxylation of alcohols and phenols
- L6 ANSWER 11 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI N-phenylcarbamate ester oligomers
- L6 ANSWER 12 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Methanetrisulfonic acid derivatives
- L6 ANSWER 13 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI 2-Hydroxyacetophenone via Fries rearrangement and related reactions. A comparative applied study
- L6 ANSWER 14 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Tris(fluorosulfonyl)methane, HC(SO2F)3
- L6 ANSWER 15 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI New electrolytes for direct methane fuel cells
- L6 ANSWER 16 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Solid catalysts for heterogeneous reactions
- L6 ANSWER 17 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Alkylation of phenols
- L6 ANSWER 18 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Esterification catalysts
- L6 ANSWER 19 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Acid-base equilibria in glacial acetic acid
- L6 ANSWER 20 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Reaction of acetylene and acetic acid. Societe des usines chimiques Rhone-Poulenc
- L6 ANSWER 21 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Reaction of oleum with AcOH or Ac20
- L6 ANSWER 22 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI The chlorination of methanetrisulfonic acid
- L6 ANSWER 23 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI The salts of methanetrisulfonic acid
- L6 ANSWER 24 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Methanetrisulfonic acid
- L6 ANSWER 25 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN

L6 ANSWER 2 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN

TI Process for the manufacture of tocyl and tocopheryl acylates

AN 2004:965239 CAPLUS

DN 141:395687

TI Process for the manufacture of tocyl and tocopheryl acylates

IN Bonrath, Werner; Haas, Alois; Hoppmann, Simone; Netscher, Thomas; Pauling, Horst

PA DSM IP Assets B.V., Neth.

SO PCT Int. Appl., 15 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

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	PAT	CENT 1	NO.			KIN	D	DATE			APPL	ICAT.	ION I	NO.		D	ATE	
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PI	WO	2004096790			A1	Al 20041111			WO 2004-EP4144						20040419			
		W:	ΑE,	AG,	AL,	AM,	AT,	AU,	AZ,	BA,	BB,	BG,	BR,	BW,	BY,	ΒZ,	CA,	CH,
			CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,
			GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KΕ,	KG,	ΚP,	KR,	ΚZ,	LC,
			LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NA,	NI,
			NO,	NZ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	.SD,	SE,	SG,	SK,	SL,	SY,
			ТJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	zw
		RW:	BW,	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	ΑZ,
			BY,	KG,	ΚZ,	MD,	RU,	TJ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,
			ES,	FI,	FR,	GB,	GR,	HU,	ΙE,	IT,	LU,	MC,	NL,	PL,	PT,	RO,	SE,	SI,
			SK,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,
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EP 2003-9522 A 20030428

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OS CASREACT 141:395687; MARPAT 141:395687

GI

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AB A process for the manufacture of tocyl acylate I [R = acyl; R1 = R2 = R3 = H] or a tocopheryl acylate I [R = acyl; R5 = R7 = R8 = Me, R5 = H, R7 = R8 = Me, etc.] comprised reacting a corresponding tocol or tocopherol with an acylating agent in the presence of a catalyst of the general formula HCR1R2R3 [wherein R1, R2 and R3 each signify the sulfo group, or R1, R2 and R3 each signify a perfluoroalkylsulfonyl group whereby at least two of R1, R2 and R3 are identical such perfluoroalkyl-sulfonyl groups, or R1 signifies the pentafluorophenyl-sulfonyl group and R2 and R3 each signify an identical perfluoroalkylsulfonyl group]. The main com. form of vitamin E, being (all-rac)- α -tocopheryl acetate I [R = acetyl; R5 = R7 = R8 = Me], can be manufactured by acylation of (all-rac)- α -tocopherol according to this process.

RE.CNT 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 5 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN TI Manufacture of α -tocopherol from the reaction of

trimethylhydroquinone with isophytol or phytol in the presence of methane trisulfonate

AN 2004:453200 CAPLUS

DN 141:23750

TI Manufacture of α -tocopherol from the reaction of trimethylhydroquinone with isophytol or phytol in the presence of methane trisulfonate

IN Bonrath, Werner; Hoppmann, Simone; Haas, Alois; Netscher, Thomas; Pauling, Horst

PA DSM IP Assets B.V., Neth.

SO PCT Int. Appl., 13 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN. CNT 1

PAIN.																		
									APPLICATION NO.									
ΡI												2003-					0030	930
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			TN,	TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC	, VN,	YU,	ZA,	ZM,	ZW		
		RW:	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ	, TZ,	ŪĠ,	ZM,	ZW,	AM,	AZ,	BY,
			KG,	KZ,	MD,	RU,	TJ,	TM,	AT,	BE,	BG	, CH,	CY,	CZ,	DE,	DK,	EE,	ES,
			FI,	FR,	GB,	GR,	HU,	IE,	IT,	LU,	MC	, NL,	PT,	RO,	SE,	SI,	SK,	TR,
			BF,	ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ	, GW,	ML,	MR,	NE,	SN,	TD,	TG
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	ΑU	2003	2702	95		A1 20040615				AU 2003-270295						20030930		
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												2003-					0030	930
	DE	1039	3642			Т5		2005	1110]	DE	2003-	1039	3642		2	0030	930
]	EP	2002-	2599	0	1	A 2	0021	121
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	CN	1701	065			Α		2005	1123			2003-				_	0030	
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		2006				A1			0126	1	US	2005-	5356	03		2	0050	519
	US	7153	984			B2		2006	1226					_				
												2002-		-	-			
										į	WO	2003-	EP10	837	I	v 2	0030	930

OS CASREACT 141:23750

AB (all-rac)- α -tocopherol is prepared by the acid-catalyzed reaction of trimethylhydroquinone with isophytol or phytol in the presence of methane trisulfonate as the catalyst in an organic solvent.

- L6 ANSWER 16 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Solid catalysts for heterogeneous reactions

AN 1975:64946 CAPLUS

DN 82:64946

TI Solid catalysts for heterogeneous reactions

IN Rona, Peter

PA IMI (TAMI) Institute for Research and Development

SO Ger. Offen., 21 pp. CODEN: GWXXBX

DT Patent

LA German

FAN.CNT 1

	PATENT NO.	KIND .	DATE	APPLICATION NO.		DATE
					-	
PI	DE 2401958	A1	19740718	DE 1974-2401958		19740116
				IL 1973-41330	Α	19730117
	US 3920582	A	19751118	US 1974-430804		19740104
	•			IL 1973-41330	Α	19730117
	GB 1446964	A	19760818	GB 1974-1839		19740115

JP 50046587 A 19750425 JP 1974-7615 19740117
IL 1973-41330 A 19730117
IL 1973-41330 A 19730117

AB Catalysts for heterogeneously catalyzed reactions were prepared by impregnation of carriers with sulfonic acids. Thus, 50 g SiO2-Al2O3 pellets were treated for 30 min with 14 g benzene-1,3-disulfonic acid in H2O at 80°, dried for 6 hr at 150°, and calcined for 6 hr at 200° to give 60 g catalyst. A H2O-C2H4 mixture of mol. ratio 1:1 was passed over this catalyst at 195° to give a C2H4-C2H5OH conversion of 0.3-0.5 mole % without splitting off acid from this catalyst.

L6 ANSWER 18 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN

TI Esterification catalysts

AN 1963:14557 CAPLUS

DN 58:14557

OREF 58:2371g-h

TI Esterification catalysts

IN Touey, George P.; Goins, Rex H.

PA Eastman Kodak Co.

SO 3 pp.

PT

DT Patent

LA Unavailable

PATENT NO. KIND DATE APPLICATION NO. DATE
US 3053884 19620911 US 1959-845336 19591009

AB CH2(SO3H)2 and CH(SO3H)3 are superior catalysts for preparing esters by treating saturated aliphatic mono- and polyhydroxy alcs. with phenyl dicarboxylic acids or saturated aliphatic carboxylic acids and their anhydrides. A lower concentration of catalyst is required and the ester produced

is nearly colorless and is heat stable. Two moles phthalic anhydride and five moles BuOH were refluxed 7 hrs. in the presence of various acid catalysts. The catalyst used, the catalyst concentration based on the phthalic anhydride, and the percent phthalic acid in the product are: CH2(SO3H)2, 0.1, 0.02; CH(SO3H)3, 0.1, 0.03; H2SO4, 0.1, 0.35; MeSO3H, 0.2, 1.6; MeC6H4SO3H, 1.0, 2.0; (CH2SO3H)2, 0.2, 0.85. Data are given which show the superiority of these two catalyst for the esterification of n-octyl alc. with adipic acid and glycerol with 2-ethylhexanoic acid.

L6 ANSWER 19 OF 25 CAPLUS COPYRIGHT 2007 ACS on STN

TI Acid-base equilibria in glacial acetic acid

AN 1953:70596 CAPLUS

DN 47:70596

OREF 47:11919f-i

TI Acid-base equilibria in glacial acetic acid

AU Smith, Thor L.; Elliott, John H.

CS Hercules Powder Co., Wilmington, DE

SO Journal of the American Chemical Society (1953), 75, 3566-71 CODEN: JACSAT; ISSN: 0002-7863

DT Journal

LA Unavailable

Values of Ho for dilute solns. $(5+10-4\ \text{to}\ 5+10-3\ \text{M})$ of 11 strong acids in AcOH containing 0.12% water were measured by use of indicators α -naphtholbenzein (I) and o-nitroaniline. Ho = -log(BH+)/(B) + pKa, where (BH+) and (B) are the concns. of the acidic and basic forms of an indicator, and pKa is the thermodynamic dissociation constant for the conjugate acid of the indicator. The pKa for I was evaluated as 0.53. The order of increasing acid strength at equal molarities is: HCl, methanesulfonic, sulfuric, carboxymethanesulfonic, chloromethanesulfonic, chlorocarboxymethanesulfonic, HBr, perchloric, methanedisulfonic, chloromethanedisulfonic, and methanetrisulfonic acids. Ho values for anhydrous solns. of 4 monobasic acids at 5 + 10-3 M were measured, and from the increased acidity found, equilibrium consts. for the reaction of the acids with water were calculated H2SO4 was found to be monobasic.

consts., Kc, of HCl, HBr, HClO4, and H2SO4 in AcOH (calculated from conductivity data

of Kolthoff and Willman (C.A. 28, 3644.1)) are 5.1 + 10-10, 1.9

+ 10-7, 9 + 10-7, and 7.4 + 10-9, resp. The fact that

values of ΔpKc from conductivity and from Ho data are in reasonable agreement shows that equilibrium in AcOH involve, primarily, undissocd.

species.

=> logoff hold

COST IN U.S. DOLLARS SINCE FILE TOTAL

FULL ESTIMATED COST ENTRY SESSION 39.10 62.28

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL

CA SUBSCRIBER PRICE ENTRY SESSION -3.90 -3.90

SESSION WILL BE HELD FOR 120 MINUTES
STN INTERNATIONAL SESSION SUSPENDED AT 06:04:45 ON 12 JUN 2007

Connecting via Winsock to STN

Welcome to STN International! Enter x:x

LOGINID: SSSPTA1623PAZ

PASSWORD:

* * * * * RECONNECTED TO STN INTERNATIONAL * * * * * * SESSION RESUMED IN FILE 'CAPLUS' AT 07:01:07 ON 12 JUN 2007 FILE 'CAPLUS' ENTERED AT 07:01:07 ON 12 JUN 2007 COPYRIGHT (C) 2007 AMERICAN CHEMICAL SOCIETY (ACS)

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 39.10 62.28

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE TOTAL ENTRY SESSION

CA SUBSCRIBER PRICE ENTRY SESSION -3.90 -3.90

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(FILE 'HOME' ENTERED AT 05:41:16 ON 12 JUN 2007)

FILE 'REGISTRY' ENTERED AT 05:41:32 ON 12 JUN 2007 E KETOISOPHORONE/CN

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FILE 'CAPLUS' ENTERED AT 05:42:06 ON 12 JUN 2007

FILE 'REGISTRY' ENTERED AT 05:42:15 ON 12 JUN 2007

E METRHANETRISULFONIC ACID/CN

E METHANETRISULFONIC ACID/CN

L2 1 E3

E SULFONIC ACID/CN

E METHANESULFONIC ACID/CN

L3 1 E3

FILE 'CAPLUS' ENTERED AT 05:44:02 ON 12 JUN 2007

L4 0 L1 AND L2

L5 396 L1

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25 L2
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           5760 L3
L7
=> toluenesulfonic
         20641 TOLUENESULFONIC
             1 TOLUENESULFONICS
         20642 TOLUENESULFONIC
                  (TOLUENESULFONIC OR TOLUENESULFONICS)
=> methanetrisulfonic
            37 METHANETRISULFONIC
=> 18(1)19
L10
             1 L8(L)L9
=> d l10 1 ti fbib abs
     Alkylation of phenols
TI
AN
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L10 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2007 ACS on STN

1963:468923 CAPLUS

DN 59:68923 OREF 59:12707d-f

TI Alkylation of phenols

McConnell, Wayne V.; Davis, Herman E. IN

PA Eastman Kodak Co.

SO 2 pp.

DT Patent

T.A Unavailable

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 3082258		19630319	US 1960-28557	19600512
				US	19600512

The preparation of 2,6-di-tert-butyl-4-methylphenol (I) from 4-methylphenol AB (II) and isobutylene using hydrated methanedi- or trisulfonic acid catalysis was described. I was useful as an antioxidant and stabilizer for fats and oils. Thus, 112 g. isobutylene was bubbled through a flask containing 108 g. II and 1.1 g. methanedisulfonic acid dihydrate (III) in 100 cc. benzene. In the initial stages the temperature varied from 25-40° due to the cooling effect of isobutylene refluxing in a dry ice-acetone cooled condenser. Thereafter the temperature was held at 40° for a total reaction time of 6 hrs. The supernatant liquid was decanted from the catalyst. Unreacted II (6%) and 2-tert-butyl-4-methylphenol (31%) conversion) were extracted with aqueous NaOH. After removal of C6H6, I was obtained (63% conversion), m. 68-9° (50% aqueous MeOH). Under the same conditions, 5.5 q. III gave an 88% conversion to I. Only a 20% conversion resulted from use of 1,2-ethanedisulfonic acid. Benzenedisulfonic acid caused polymerization of isobutylene. When p-toluenesulfonic acid or H2SO4 was used in concentration of 5% based on the weight of II the product

had poorer color and odor. White, odorless I could also be prepared in 84 and 80% conversions, resp., using 2.2 g. III and no solvent or using 1% by weight methanetrisulfonic acid trihydrate.

=> logoff hold SINCE FILE COST IN U.S. DOLLARS TOTAL ENTRY SESSION 62.40 FULL ESTIMATED COST 85.58 SINCE FILE DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) TOTAL. ENTRY SESSION CA SUBSCRIBER PRICE -4.68 -4.68 SESSION WILL BE HELD FOR 120 MINUTES
STN INTERNATIONAL SESSION SUSPENDED AT 07:22:07 ON 12 JUN 2007